

*Free Home Sample Collection 9999 778 778

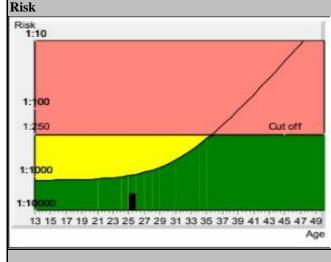


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Date of Report 07-12-18
PRISCA 5.0.2.37

					TRISCA	5.0.2.57
Patient Data						
Name		M	rs Chinnari	Patient ID		011812060069
Birthday			07-07-93	Sample ID		10352116
Age at delivery			25.4	Sample Date		06/12/2018
Gestational age			13+3			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes		no	Pregnancies	
Smoker	no	Origin		Asian		
Biochemical Data				 Ultrasound D	ata	

Biocnemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	13+3	
PAPP-A	3.55 mIU/ml	0.65	Method	CRL (<>Hadlock)	
fb-hCG	54.51 ng/ml	1.33	Scan Date	06-12-18	
Risks at sampling date			CRL	71.9	
Age Risk		1:972	Nuchal translucency MoM	0.56	
Biochemical T21 Risk		1:1159	Nasal Bone	Present	
Combined Trisomy 21 Risl	ζ	1:6720	Sonographer	DR.SAHIL LOOMBA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS,DNB	
			D 1 G 1 D11 G	4. G	



Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 6720 women with the same data, there is one woman with a trisomy 21 pregnancy and 6719 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines. The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!